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October 10, 1991

Ms. Marcia Bailey U.S. Environmental Protection Agency Region 10 1200 Sixth Avenue Seattle, WA 98101

Subject:

Contract No. 68-W9-0009, Work Assignment No. 12R10047

Operation and Maintenance Inspection

Yakima Agricultural Research Laboratory, Washington

Dear Ms. Bailey:

PRC Environmental Management, Inc. (PRC) is pleased to submit one copy of the Operation and Maintenance Inspection Report for the Yakima Agricultural Research Laboratory (YARL). Also, analytical data from the YARL inspection has been entered into the RCRA Groundwater Database as required by the Regional Groundwater Management Order (R107500.1). Enclosed is a computer disk and a hard copy of the data. Please forward the database information to Matt Gubitosa.

If you have any questions or comments, please contact Ben Farrell or me at 624-2692.

Sincerely,

Gary A. Bruno

Environmental Geologist

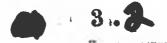
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Enclosures

cc: Vicky Tapang, EPA Region 10 (without enclosures)

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U.S. Environmental Protection Agency
Office of Waste Programs Enforcement
Contract No 68-W9-0009

YAKIMA AGRICULTURAL RESEARCH LABORATORY

YAKIMA, WASHINGTON

OPERATION AND MAINTENANCE INSPECTION REPORT

TES 12

Technical Enforcement Support at Hazardous Waste Sites Zone IV Regions 8, 9, and 10

prc

PRC Environmental Management, Inc.

draft

YAKIMA AGRICULTURAL RESEARCH LABORATORY

YAKIMA, WASHINGTON

OPERATION AND MAINTENANCE INSPECTION REPORT

Prepared For

U.S. ENVIRONMENTAL PROTECTION AGENCY Office of Waste Programs Enforcement Washington, D.C. 20460

Work Assignment No. 12R10047

EPA Region 10

Telephone No.

 EPA ID. No.
 WAD 120513957

 Date Prepared
 October 10, 1991

 Contract No.
 68-W9-0009

 Work Assignment No.
 212-R1004709

Prepared by

PRC Environmental

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206/553-0684

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EXECUTIVE SUMMARY

PRC Environmental Management, Inc. (PRC), received Work Assignment No. 112R10047 from the U.S. Environmental Protection Agency (EPA) to conduct an operation and maintenance (O&M) inspection at the Yakima Agricultural Research Laboratory (YARL) in Yakima, Washington. At the request of EPA, PRC performed the O&M inspection to evaluate how the facility operates and maintains its groundwater monitoring system in terms of pertinent RCRA regulations and permit requirements.

and the sampling methods used by the facility contractor. In general, the monitoring wells maintenance and construction, and sampling procedures used by the facility contractor are adequate to ensure the collection of representative groundwater samples. In accordance with 40 CFR 265.92, the sampling and analysis plan (Hong West and Associates 1990a) provides adequate procedures and techniques for groundwater sample collection, sample preservation and shipment, analytical procedures, and chain-of-custody control.

PRC makes the following recommendations regarding field sampling procedures observed during the site visit:

- The pump rate observed during the collection of volatile organic compounds was commonly too high. The facility contractor should reduce the pump rate to minimize the potential for sample volatilization.
- The teflon tubing connected to the pump discharge outlet was too short at monitoring well MW-D. Because of this situation, potentially contaminated groundwater was spilled on the ground surface, and the lip of sample containers was allowed to contact the teflon tubing. A longer length of tubing should be installed.
- Purge water collected in drums should be covered until chemical analysis has been completed.
- The facility contractor should have extra glassware available during sampling.
- Total well depth was not measured during the sampling event. The sampling and analysis plan states that total well depth would be monitored on a monthly basis. In accordance with EPA (1986a), total well depth should be measured regularly in order to monitor the wells for siltation problems and well integrity.

The facility contractor found no pesticide or volatile organic contamination during the May 6, 1991 Round 9 of groundwater sampling. Metals such as calcium, sodium, magnesium, and potassium were detected by the facility contractor in relatively high concentrations in both upgradient and downgradient wells. These relatively high background concentrations

could be related to local agricultural practices. A concentration of 3.5 micrograms per liter $(\mu g/L)$ lead was detected in one of the downgradient monitoring wells, MW-A.

Results from split samples collected by PRC show low concentrations of lead and arsenic in groundwater samples from both upgradient and downgradient wells at the YARL site. The arsenic was detected in concentrations below the quantitation limits of the facility laboratory. Concentrations of metals such calcium, sodium, magnesium, and potassium were detected at concentrations comparable to those reported by the facility contractor. No significant differences were found between dissolved and total metals analysis in the split samples collected by PRC, indicating that most of the metals are dissolved in the groundwater. Split sample results also show low concentrations of chloroform and endosulfan sulfate in all of the downgradient monitoring wells, but not in upgradient monitoring well MW-D. However, both of the contaminants were found in concentrations below the quantitation limit of the facility laboratory.

The good comparability between split sample data and the facility analytical data suggests that the facility analytical program is adequate. The detection of contaminants below the facility laboratory quantitation limits by the EPA approved laboratories shows the potential for very low levels of groundwater contamination to exist and not be detected by the facility laboratory.

The EPA approved closure plan states that the facility will completely characterize the YARL site hydrogeology and assess the interconnection of the upper and lower aquifers at the site. PRC notes the following deficiencies regarding the YARL hydrogeologic site characterization:

- Anomalous water level measurements at monitoring well MW-B should be explained rather than disregarded.
- The cause for the monthly variations in the direction of groundwater flow should be investigated. The facility should evaluate the seasonal use of nearby irrigation wells and ditches to determine whether irrigation practices are affecting groundwater flow directions at the site.
- The degree of interconnection between the upper alluvial aquifer and underlying aquifers has not been established. At a minimum, the facility should define and more completely characterize the underlying aquifers through examination of well logs or permits for local water supply wells and discussion with parties involved in local groundwater monitoring or supply of groundwater.

1.0 INTRODUCTION

This report describes the operation and maintenance (O&M) inspection conducted on May 6, 1991 at the Yakima Agricultural Research Laboratory (YARL) site in Yakima, Washington.

The specific objectives of the YARL O&M inspection were to:

- Evaluate the compliance of the groundwater monitoring system with the Resource Conservation Recovery Act (RCRA) interim status groundwater monitoring regulations (40 CFR 265 Subpart F) and protocols specified in EPA (1986)
- Determine whether sampling devices are in working order and are properly maintained
- Evaluate the facility sampling and analysis plan
- Determine whether individual monitoring wells yield representative groundwater data
- Evaluate groundwater contamination from analytical results of split groundwater samples received by PRC
- Evaluate the analytical program of the facility through the comparison of facility and split sample analytical results

PRC personnel present onsite were Ben Farrell, geologist, and Julie Howe, environmental scientist. Technical points were discussed with Sweet-Edwards/EMCON, Inc. geologist Becky Hylland. A photographic log of the site visit is shown in Appendix A.

2.0 SETTING AND SITE HISTORY

The YARL site is a RCRA-regulated land disposal facility located at 3706 West Nob Hill Boulevard in Yakima, Washington. The 9.5 acre site is situated in a residential area as shown in Figure 1. Three schools, two hospitals, and three shopping centers are located within one-half mile of the site (Tetra Tech 1989). The objective of this inspection was to evaluate the groundwater monitoring system associated with a septic tank/drainfield system.

The laboratory is administered by the U.S. Department of Agriculture. Originally an orchard, the YARL site began to be used for pesticide research in 1961. Several types of pesticide wastes and solvents were disposed of directly on the ground until 1965. Between 1965 and 1985, wastes were discharged to a septic tank and drainfield via a sink and/or a concrete pad at the facility (Tetra Tech 1989).

The septic system, which is in the process of removal as part of the facility closure effort, consisted of a 300-gallon concrete tank connected to a 4-inch diameter, 30-foot long drain tile. The drain tile sloped to the southeast and was installed between 2 and 4 feet below ground surface (bgs). Overflow effluent from the tank was discharged through the drain tile. Beginning in approximately 1965, pesticide-contaminated laboratory equipment was rinsed in a sink in the pesticide storage building, and the rinse water drained from the sink into the septic tank.

In approximately 1974, a 165-foot square concrete pad was installed for the purpose of cleaning pesticide application and miscellaneous farm equipment. A surface drain was constructed along the perimeter of the concrete pad. This surface drain was connected to the septic tank with a 4-inch concrete pipe. Field sprayers, tractors, and other equipment were rinsed on this pad.

Disposal of pesticide products via the lab sink stopped by 1984 and the surface drain surrounding the concrete wash pad was sealed in June 1985. Roughly 5,000 gallons of rinsate from pesticide application equipment and a maximum of 250 gallons of various solvents and pesticide solutions were discharged through the system yearly (Biospherics 1988). The presence of highly permeable sands and gravels caused concern that pesticides and solvents had leached into the shallow drinking water aquifer (Tetra Tech 1989).

YARL submitted a RCRA Part A permit application in September 1980. A preliminary assessment and site investigation pursuant to the Comprehensive Environmental Response, Conservation, and Liability Act (CERCLA) was conducted in June 1982. The site was proposed for the Superfund National Priority List in December 1982. YARL is currently ranked 1026 among the 1073 sites on the National Priorities List (56 Federal Register 5605, February 1991).

A closure plan for the septic tank and drainfield system that includes a monitoring plan for sampling and analyzing groundwater and soils was submitted by YARL in January 1985. In March 1987, YARL submitted a revised version of this closure plan, which was approved by Washington Department of Ecology in May 1987. However, in September 1987, EPA determined that the closure plan did not meet the requirements of 40 CFR 265 Subpart G and requested that a revised closure plan be submitted to EPA after implementation of a groundwater monitoring system pursuant to 40 CFR 265 Subpart F. This groundwater monitoring system, consisting of four wells, was installed in April 1988. A revised closure plan was submitted and subsequently approved by EPA on January 30, 1990. The central component of the EPA-approved plan is to achieve clean closure under 40 CFR 265 subpart G. As required by the approved closure plan, three additional monitoring wells were drilled and completed by July 1990. The purpose of the three additional wells was to determine the vertical hydraulic gradient at one locality, and to

provide a more complete monitoring network, which was deemed necessary due to the variation in groundwater flow directions at the site (Tetra Tech 1989).

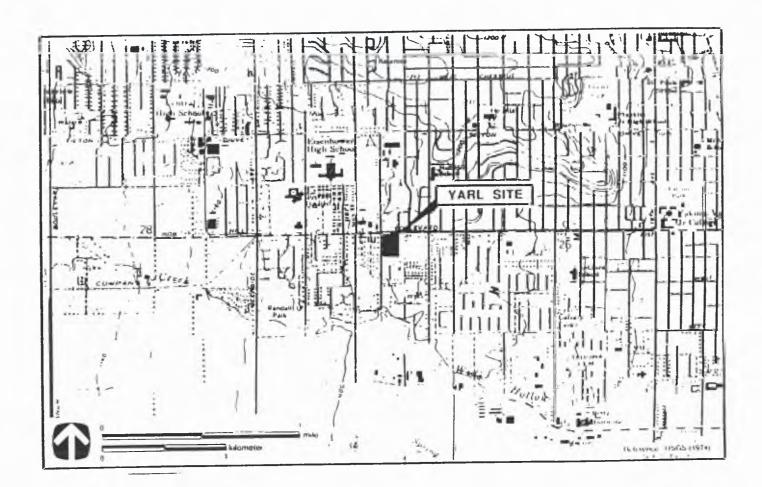
3.0 SITE GEOLOGY AND HYDROGEOLOGY

There are two main aquifers of concern underlying the YARL site. The shallow aquifer consists of alluvial sands and gravels of the Ellensburg Formation. The deeper aquifer is located in interflow zones of the underlying Columbia River basalts (Tetra Tech 1989). The total thickness of the Ellensburg Formation at YARL has not been determined. Basalt bedrock was not encountered at a maximum drilling depth of 123 feet. The water table at the site is located 34 to 38 feet bgs. The direction of groundwater flow in the upper aquifer is generally to the south-southeast toward Wide Hollow Creek (Tetra Tech 1989). The direction of groundwater flow in the upper aquifer may vary by as much as 45 degrees between consecutive months (Tetra Tech 1989).

4.0 GROUNDWATER MONITORING SYSTEM

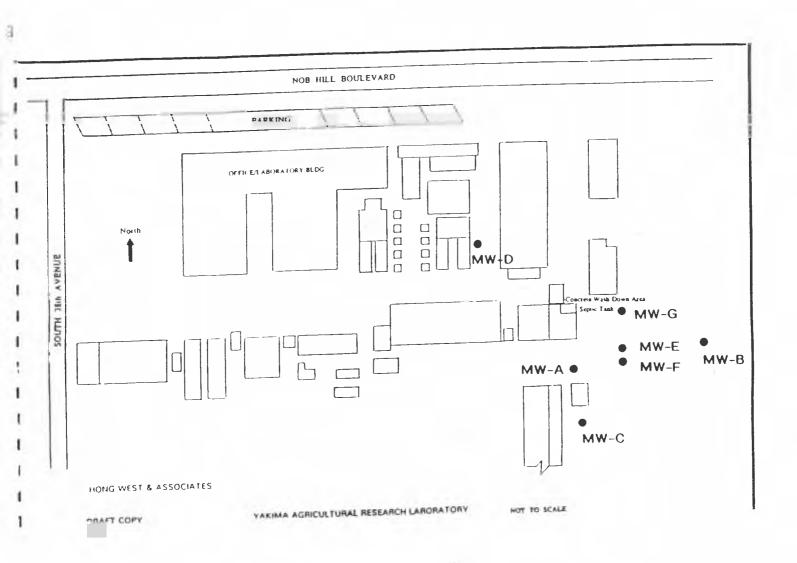
The YARL groundwater monitoring system consists of seven wells (MW-A through MW-G). Monitoring well locations are shown in Figure 2. Monitoring well MW-D is located upgradient, while all of the other wells are either cross-gradient or downgradient of the septic tank and drainfield, depending on variations in the direction of groundwater flow (Tetra Tech 1989). Monitoring well MW-E was installed as a piezometer at a depth of 125 feet bgs near the bottom of the shallow alluvial aquifer in order to generate information regarding vertical hydraulic and chemical gradients within the upper aquifer (Hong West and Associates, 1990a). All of the other wells are screened in uppermost 10 feet of the shallow aquifer.

FIGURE 1 LOCATION MAP OF YAKIMA AGRICULTURAL RESEARCH LABORATORY



From Tetra Tech (1989)

FIGURE 2 YARL WELL LOCATION MAP



5.0 SITE INSPECTION

On May 6, 1991, Ben Farrell, geologist, and Julie Howe, environmental scientist, conducted an operation and maintenance inspection at YARL. The inspection corresponded to the fourth quarterly groundwater sampling event as described in the facility project plan (Hong West and Associates 1990a). The weather was generally sunny with temperatures ranging between 60-70°F. PRC personnel observed sampling procedures at the following wells: MW-A, MW-B, MW-C, MW-D, MW-E, MW-F, and MW-G. Split-groundwater samples were received by PRC personnel from monitoring wells MW-D, MW-G, MW-A, MW-F, and MW-E. A photographic log of the site visit is presented in Appendix A Sweet-Edwards/EMCON field notes, PRC field notes, an operation and maintenance inspection checklist, a YARL analytical data summary, a PRC analytical data summary, and potentiometric surface maps are presented in Appendices B through G, respectively.

5.1 WELL MAINTENANCE AND ABOVE-GROUND WELL CONSTRUCTION

The above-ground portion of the monitoring wells appeared to be adequately maintained. Monitoring well MW-D was constructed to grade because of heavy traffic in that portion of the YARL site. The lid to the protective casing was secured and required an allen wrench to be opened, but was not locked. Rust visible on the well cap indicated the probability of moisture collecting inside the well vault. With the exception of monitoring well MW-D, all of the other inspected wells were securely locked. The wells constructed above the ground were surrounded by a three protective posts approximately 3 feet high, and had lockable protective outer casings. Each well was surveyed at a marked point on the top rim of the outer well casing. None of the inspected wells were labelled. The protective outer casing was set into a triangular concrete pad. In all cases, the concrete pad was slightly raised above the ground surface and fit tightly against the protective outer casing. The top of the polyvinyl chloride (PVC) inner casing was covered by a well cap attachment used for the connection of the dedicated pump and automatic controller.

5.2 GROUNDWATER PUMPS

A dedicated well wizardtm bladder pump was used to sample all wells. The pump was connected to a model 3013 automatic controller that supplied pressurized nitrogen as the pump gas. The pressurized nitrogen was used to ensure that no ambient air contaminated the groundwater samples.

5.3 WATER-LEVEL MEASUREMENTS

Water levels were measured to the nearest 0.01 foot at all of the monitoring wells on the same day prior to sampling any of the wells. These measurements were not repeated after purging and sampling. Depth to water was measured with a Slope Indicator Inc. electronic probe. Total well depth was not measured. In accordance with the sampling and analysis plan, the well probe was decontaminated by rinsing the coiled tape and the probe tip with methanol and then twice with distilled water.

5.4 WELL FURGING AND ENVIDONMENTAL PARAMETERS

The monitoring wells were purged and sampled in the following order: MW-D, MW-G, MW-B, MW-A, MW-F, MW-E, and MW-C. As specified in the project sampling and analysis plan, the above order of sample collection is basically from upgradient to downgradient wells. Monitoring well MW-C was sampled last because of its prior history of volatile organic contamination.

The sampling and analysis plan stipulates that a minimum of five well casing volumes will be purged before the commencement of sampling. With the exception of monitoring well MW-G, three well casing volumes were purged at all of the inspected wells. After each well casing volume was discharged, a groundwater sample was analyzed for conductivity, pH, and temperature. Measurements of the conductivity and pH were made using a DSPH3 meter and temperature measurements were made using a Taylor model 21431 thermometer. The conductivity and pH meter was calibrated before sample collection was initiated at monitoring well MW-D. Results were tabulated on field data sheets (Appendix B). The sampling and analysis plan states that well purging will continue even beyond five well casing volumes until successive measurements of the field parameters fall within 10 percent. In general, the field groundwater parameters for the inspected wells stabilized by the collection of the third well casing volume. An exception to this trend occurred at monitoring well MW-G where four well casing volumes were collected. Well volumes were calculated on the field data sheets (Appendix B).

Purged groundwater was pumped into buckets of known volume and transferred into a 55 gallon drum for storage until the calculated purge volume was reached. The containerized purge water was left open on the facility premises to evaporate in the sun. It is possible that during the rainy season, the filled 55-gallon drums could overflow onto the ground surface.

SAMPLE COLLECTION

5.5

After well purging, groundwater samples were collected by the facility contractor using clean gloves. Split groundwater samples were collected for the following parameters and analyzed using methods specified in EPA (1986b):

- volatile organic compounds (SW-846 8240)
- organo-chlorine pesticides (SW-846 8080)
- chlorinated herbicides (SW-846 8150)
- organo-phosphorus pesticides (SW-846 8140)
- total metals (SW 846 methods 6010, 7470)

In accordance with the sampling and analysis plan, the facility collected a sample duplicate at monitoring well MW-D and a trip blank. All containers used by the facility contractor for the collection of volatile organics and metals were preserved by the laboratory prior to the sampling event. Volatile organic samples were preserved with concentrated hydrochloric acid, and metal samples were preserved with nitric acid.

Samples were collected in the following order: volatile organic compounds; pesticides and herbicides; and metals. The following sample containers were used for sample collection:

volatile organic compounds
pesticides and herbicides
total metals

2 x 40 mL glass vial
1 x 4 L amber glass jug
1 x 1 L polyethylene

The sampling and analysis plan specifies that all samples collected for metals analysis would be analyzed for filtered total metals rather than total metals. Results reported by the facility laboratory were for total metals only.

At monitoring well MW-D, the plastic pump outlet tube was too short to allow for the proper filling to the bucket used for purging and the large 4 liter containers used for pesticide samples. This situation caused potentially contaminated groundwater to be spilled on the ground. Also, during the collection of samples at this well, the sample container lips were periodically in contact with the pump outlet tube. In order to prevent container contamination, the sample containers should not come in contact with the pump discharge tube. A longer pump outlet tube should be installed at monitoring well MW-D.

During the collection of volatile organic samples, the pump rate was occasionally too high for the collection of representative samples. EPA (1986a) states that pump rates during volatile organic sample collection should not exceed 100 milliliter per minute (mL/minute). The observed

pump rate was considerably in excess of 100 mL/minute. The groundwater sampling stream was often quite aerated and preservatives were occasionally flushed from the sample containers. According to the facility contractor, the flow throttle on the automatic controller was not functioning properly. The facility contractor tried to control the flow rate by partially closing the valve to the pressurized nitrogen canister and by adjusting the refill and discharge controls on the automatic controller.

During the collection of volatile organic analysis (VOA) vials, the septa fell out of VOA vials repeatedly during sampling. Because no extra VOA vials were brought along by the facility contractor, the septa were decommunicated using distilled water and were then used for the collection of samples.

Sample containers were labelled in the field prior to sampling. After sampling, the containers were placed in coolers with ice. Discussions with the facility contractor's personnel revealed that after sample collection, a chain-of-custody form is placed in each cooler and the cooler is sealed with chain-of-custody seals. Samples were shipped the same day via Federal Express to Biospherics Inc. in Beltsville, Maryland for chemical analysis.

6.0 FACILITY ANALYTICAL DATA

Facility analytical data for the May 6, 1991 sampling event is summarized in Hong West and Associates (1991c). The data summary table from the above report is included as Appendix E. Several metal compounds were detected above quantitation limits. The concentrations of calcium, sodium, magnesium, and potassium are relatively high (NAS 1977). These metals are also present in the upgradient well and are not considered site-related contaminants. These relatively high background concentrations could be related to local agricultural practices. A concentration of $3.5 \mu g/L$ lead was detected in monitoring well MW-A. Concentrations of volatile organic compounds, herbicides, and pesticides were below the laboratory quantitation limits.

A The table presented in Appendix F analytical results were submitted to EPA by PRC to EpA in July 1972 (PRC 1992),

A summary of results for the split groundwater samples received by PRC for the YARL are shown in Appendix F. In an effort to meet quality assurance/quality control objectives, PRC submitted a trip blank, and an environmental duplicate as well as a matrix spike/matrix spike duplicate for chemical analysis. Data was validated using the guidelines established by EPA (1988a,b). For a complete description of the data validation for the split groundwater samples analyzed for the YARL site see the data validation report (PRC 1991).

Split sample inorganic analytical results indicate relatively high concentrations of calcium, sodium, magnesium, and potassium similar to those reported by the facility contractor. Arsenic and lead were present at low concentrations (a maximum of 7.2 μ g/L and 17.2 μ g/L respectively) in groundwater samples from both upgradient and downgradient wells. The facility contractor did not find arsenic in concentrations above the laboratory quantitation limit. However, the laboratory quantitation limit of the facility laboratory for arsenic (10 μ g/L) is higher than the values for the split samples. The facility contractor found lead at a concentration of 3.5 μ g/L in monitoring well MW-A. Split groundwater samples from this well yielded the highest

Split sample results for volatile organic compounds showed low concentrations of methylene chloride, acetone, tetrachloroethene, and chloroform. Concentrations of all of these compounds were estimated below the practical quantitation limit and were qualified J.

Methylene chloride is a common laboratory contaminant and was only found in the trip blank.

Acetone was found at very low concentrations from two of the monitoring wells and is also a common laboratory contaminant. Tetrachloroethene was found only in very low concentrations in two of the wells. The upgradient well (MW-D) showed the highest concentration (3 \mu g/L) of this compound. Chloroform was found at very low concentrations (1-3 \mu g/L) in all of the downgradient wells, but not in the upgradient well. The distribution of chloroform in the downgradient wells at YARL suggests that it may be a site related contaminant. The concentrations of the first content and the significantly below proposed subjects after leads (55 FR 307M) and the model toxic leads of the first leads of the leads of the

For pesticides and herbicides, split sample results show the presence of endosulfan sulfate in low concentrations $(0.1-0.2~\mu g/L)$ in all of the downgradient wells, but not in the upgradient well. The distribution of endosulfan sulfate suggests that it is a site related contaminant. Analysis of split samples indicated low levels of contamination that were not reported by the facility laboratory. In all such cases, the facility laboratory did not report results below the quantitation limit. However, the EPA contract laboratory reported results below the quantitation limit and greater than the method detection limit as estimates, which are designated J. This suggests that very low levels of groundwater contamination could occur and not be detected by the analytical testing program of the facility.

8.0 HISTORICAL ANALYTICAL DATA

Data from the August 1990 round of groundwater sampling (Round 1) showed low levels of mercury contamination below the maximum contaminant level for drinking water established in 40 CFR 264.93 (Hong West and Associates 1990b). During the November 14, 1990 round of groundwater monitoring (Round 2), the following pesticide compounds were detected in

groundwater from the YARL site: malathion, heptachlor, 4,4'-DDT, and heptachlor epoxide (Hong West and Associates 1991a). None of these compounds have been detected subsequently. Analytical data collected during a September 1988 comprehensive groundwater monitoring evaluation showed detectable concentrations of chloroform (1-12 μ g/L) in downgradient wells MW-A, MW-B, and MW-C (Tetra Tech 1989). Arsenic was also detected (26 μ g/L) during the comprehensive groundwater monitoring evaluation sampling event in monitoring well MW-A (Tetra Tech 1989).

The split samples collected by PRC confirm the presence of site related chloroform in the groundwater. Though arsenic was detected at low levels in the most recent split samples and has been detected in samples from downgradient monitoring wells at the facility in the past, it is not clear that arsenic is a site related contaminant because it was also found in groundwater samples from the upgradient well.

9.0 GROUNDWATER SITE CHARACTERIZATION

The EPA approved closure plan and the project work plan states that the facility will characterize the uppermost aquifer and assess the hydraulic interconnection of the uppermost and next lower aquifer. In addition, the Code of Federal Regulations 40 CFR 265.94 (b)(2) stipulates that for facilities showing groundwater contamination the rate and extent of migration of the hazardous waste constituent in the groundwater must be documented yearly. The rate and extent of contamination cannot be documented without a complete characterization of the site hydrogeology. PRC notes the following shortcomings in the hydrogeologic site characterization.

Potentiometric surface maps have been constructed on a monthly basis since August 1990 (Appendix G). These maps show that the direction of groundwater flow is variable, ranging from nearly easterly flow to nearly southerly flow. The cause of the fluctuations in groundwater flow direction, while presumably related to irrigation practices, has not been documented. The facility should evaluate the effect of nearby irrigation wells and ditches on the direction of groundwater flow.

The potentiometric surface maps prepared by the facility contractor also show a level of certainty and detail not possible from the collected data. Where approximated, potentiometric surface lines should drawn as dashed rather than solid lines. Monitoring wells MW-E and MW-B were not used in the creation of the potentiometric surface maps. Monitoring well MW-E was legitimately excluded because of the screened interval of the well was deep in the upper aquifer. Monitoring well MW-B was excluded because water-level data from this well consistently produced anomalous flow patterns (Hong West and Associates 1991a,b,c). Monitoring well MW-

B is constructed with the well screen placed at the same depth and intersecting similar lithology as the other shallow monitoring wells, and appears to be constructed in the same manner (Biospherics 1988). Rather than excluding the anomalous water level data for monitoring well MW-B, the cause for the anomalous water level data should be determined and reported.

A consistent upward vertical hydraulic gradient has been determined by comparing water levels measured at shallow monitoring wells MW-F and the deeper piezometer MW-E. The vertical hydraulic gradient has averaged roughly .01 ft/per ft upward. An exception to this trend occurred in July of 1991 where the upward vertical hydraulic gradient was .028 ft/per ft (Hong West and Associates, 1991c). These results suggest that the well locality is within a zone of groundwater discharge. Caution should be exercised when interpreting these results as an upward vertical gradient has not been documented across the entire site.

The degree of interconnection between the shallow aquifer and deeper aquifers below the site has not been established at the site. Furthermore, the lower aquifer has not been well defined.

10.0 SUMMARY AND RECOMMENDATIONS

PRC assessed all of the monitoring wells at the YARL site and determined that the maintenance and above-ground construction of the monitoring wells, and the sampling protocols used by the facility contractor were generally adequate to ensure the collection of representative groundwater samples. In accordance with 40 CFR 265.92, the sampling and analysis plan provides adequate procedures and techniques for groundwater sample collection, sample preservation and shipment, analytical procedures, and chain-of-custody control.

PRC makes the following recommendations regarding field sampling procedures observed during the site visit:

- The pump rate observed during the collection of volatile organic compounds was commonly too high. The facility contractor should reduce the pump rate to minimize the potential for sample volatilization.
- The teflon tubing connected to the pump discharge outlet was too short at monitoring well MW-D. Because of this situation, potentially contaminated groundwater was spilled on the ground surface, and the lip of sample containers was allowed to contact the teflon tubing. A longer length of tubing should be installed.
- Purge water collected in drums should be covered until chemical analysis has been completed.

- The facility contractor should have extra glassware available during sampling.
- Total well depth was not measured during the sampling event. The sampling and analysis plan states that total well depth would be monitored on a monthly basis. In accordance with EPA (1986a), total well depth should be measured regularly in order to monitor the wells for siltation problems and well integrity.

The facility contractor found no pesticide or volatile organic contamination during the May 6, 1991 Round 9 groundwater sampling. Metals such as calcium, sodium, magnesium, and potassium were detected by the facility contractor at relatively high concentrations in both upgradient and downgradient wells. These relatively high background concentrations could be related to local agricultural practices. A concentration of 3.5 μ g/L lead was detected in one of the downgradient monitoring wells, MW-A.

Results from split samples collected by PRC show low concentrations of lead and arsenic in groundwater samples from both upgradient and downgradient wells at the YARL site. The arsenic was detected in concentrations below the quantitation limits of the facility laboratory. Concentrations of metals such calcium, sodium, magnesium, and potassium were detected in close to the same concentrations as those reported by the facility contractor. No significant differences were found between split sample dissolved and total metals analysis, indicating that most of the metals are dissolved in the groundwater. Split sample results also show low concentrations of chloroform and endosulfan sulfate in all of the downgradient monitoring wells, but not in upgradient monitoring well MW-D. However, both of the contaminants were found in concentrations below the quantitation limit of the facility laboratory.

The good comparibility between independently analyzed EPA and facility analytical data suggests that the facility analytical program is adequate. The detection of contaminants below the facility laboratory quantitation limits by the EPA approved laboratories shows the potential for very low levels of groundwater contamination to exist and not be detected by the facility laboratory.

The EPA-approved closure plan states that the facility will completely characterize the YARL site hydrogeology and assess the interconnection of the upper and lower aquifers at the site. PRC notes the following deficiencies regarding the YARL hydrogeologic site characterization:

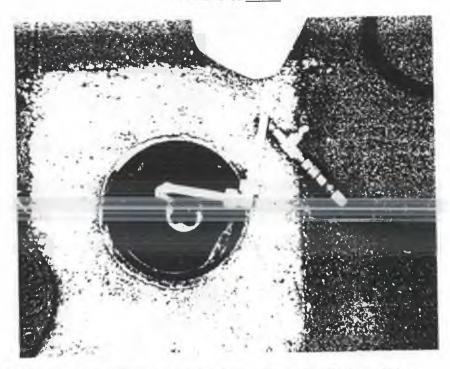
Anomalous water level measurements at monitoring well MW-B should be explained rather than disregarded.

- The cause for the monthly variations in the groundwater flow direction should be investigated. The facility should evaluate the seasonal use of nearby irrigation wells and ditches to determine whether irrigation practices are affecting groundwater flow direction at the site.
- The degree of interconnection between the upper alluvial aquifer and underlying aquifers has not been established. At a minimum, the facility should define and more completely characterize the underlying aquifers through examination of well logs or permits for local water supply wells and discussion with parties involved in groundwater monitoring or supply of groundwater locally.

11.0 REFERENCES

- Biospherics 1988. Preliminary Soil and Groundwater Investigation of the United States
 Department of Agriculture Research Laboratory in Yakima, Washington. December 23, 1988.
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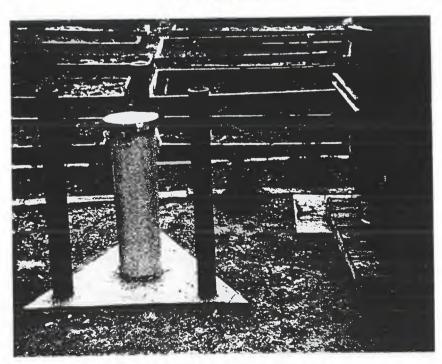
APPENDIX A PHOTOGRAPHIC LOG



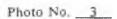
Date: May 6, 1991 Picture Taken By: Julie Howe Direction Facing: N/A

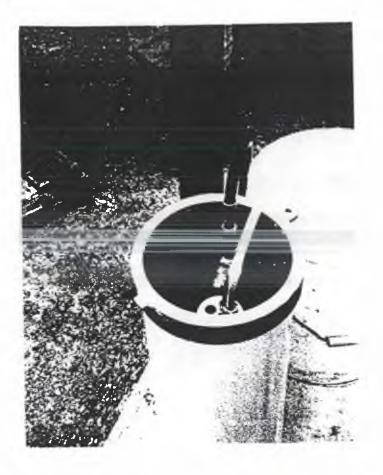
Picture Description: Monitoring well MW-D. Note flush mount, teflon tubing, and distilled water container.

Photo No. 2



Date: May 6, 1991 Picture Taken By: Julie Howe Direction Facing: East
Picture Description: Monitoring well MW-G.





Date: May 6, 1991

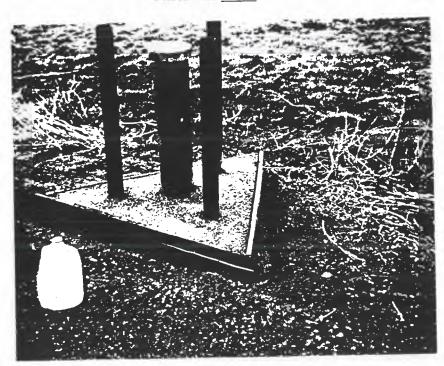
Picture Taken By: Julie Howe

Direction Facing: N/A

Picture Description Puraing at

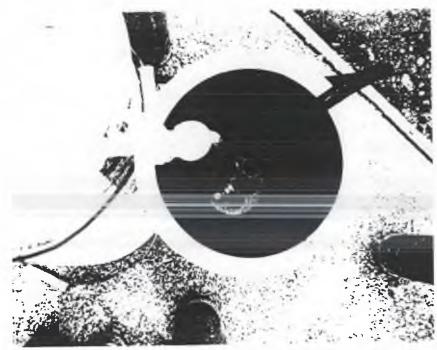
monitoring well MW-G.

Photo No. 4



Date: May 6, 1991 Picture Taken By: Julie Howe Direction Facing: N/A
Picture Description: Monitoring well MW-B.

Photo No. _5



Date: May 6, 1991 Picture Taken By: Julie Howe Direction Facing: N/A

Picture Description: Purging at monitoring well MW-B.



Photo No. 6

Date: May 6, 1991	
Picture Taken By: <u>Julie Howe</u>	
Direction Facing: South	
Picture Description: Taking	
water-level measurement at	
monitoring well MW-A.	



Date: May 6, 1991 Picture Taken By: Julie Howe Direction Facing: N/A
Picture Description: Close-up of monitoring well MW-A.

Photo No. 8

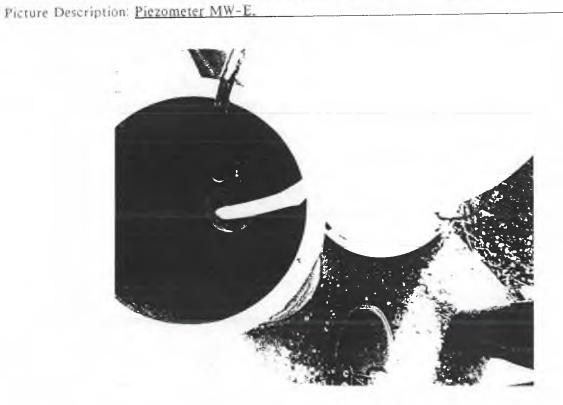


Date: May 6, 1991 Picture Taken By: Ben Farrell Direction Facing: East

Picture Description: Purging at monitoring well MW-E (left) and protective casing of monitoring well MW-F (right).



Date: May 6, 1991 Picture Taken By: Ben Farrell Direction Facing: Southeast



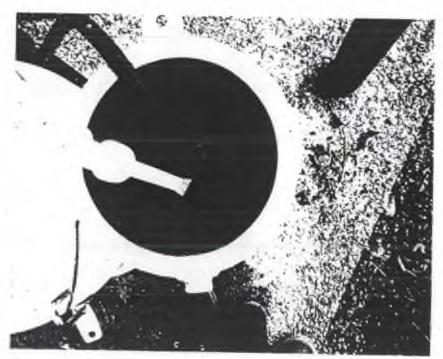
Date: May 6, 1991 Picture Taken By: Ben Farrell Direction Facing: N/A
Picture Description: Purging at piezometer MW-E.



Date: May 6, 1991 Picture Taken By: Ben Farrell Direction Facing: East

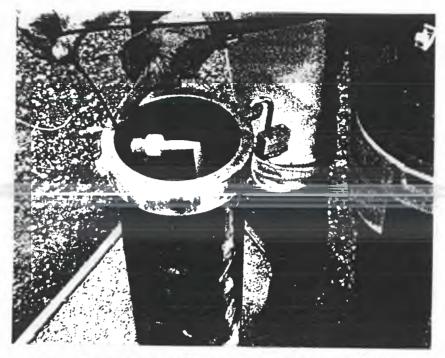
Picture Description: Monitoring well MW-C. Note pallet resting against well casing.

Photo No. 12



Date: May 6, 1991 Picture Taken By: Ben Farrell Direction Facing: N/A

Picture Description: Interior of casing at monitoring well MW-C.



Date: May 6, 1991 Picture Taken By: Julie Howe Direction Facing: N/A

Picture Description: Taking water-level measurement at monitoring well MW-B. Water level is measured at surveyed mark on casing rim as shown in the photograph.



Date: May 6, 1991 Picture Taken By: Julie Howe Direction Facing: N/A
Picture Description: Taking water-level measurement at monitoring well MW-C.



Date: May 6, 1991 Picture Taken By: Julie Howe Direction Facing: West

Picture Description: Methanol rinse of probe at monitoring well MW-E.

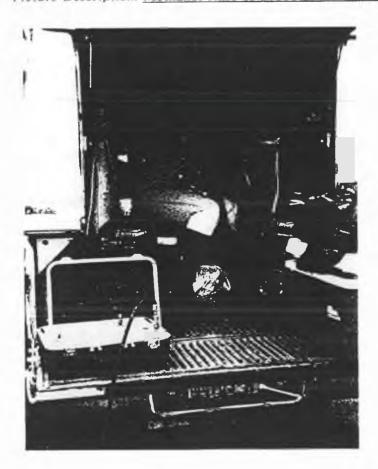


Photo No. 16

Date: May 6, 1991
Picture Taken By: Julie Howe
Direction Facing: N/A
Picture Description: Nitrogen
canisters and pump compressor.

Photo No. 17



Date: May 6, 1991 Picture Taken By: Julie Howe Direction Facing: N/A

Picture Description: Sampling for conductivity, pH, and temperature at monitoring well MW-D.



Photo No. 18

Date: May 6, 1991
Picture Taken By: Ben Farrell
Direction Facing: South
Picture Description: Purging at
monitoring well MW-C.



Date: May 6, 1991

Picture Taken By: Ben Farrell

Direction Facing: N/A

Picture Description: pH and

conductivity meter and container used

to sample environmental parameters.

Meter sensors were allowed to rest on

dirty truck bed without subsequent
decontamination.

Photo No. 20

Date: May 6, 1991

Picture Taken By: Ben Farrell

Direction Facing: West

Picture Description: Measuring pH,

temperature, and conductivity at

monitoring well MW-C.

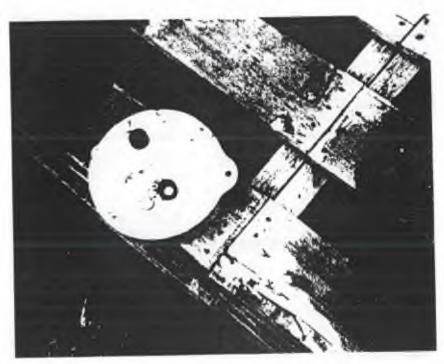


Photo No. 21



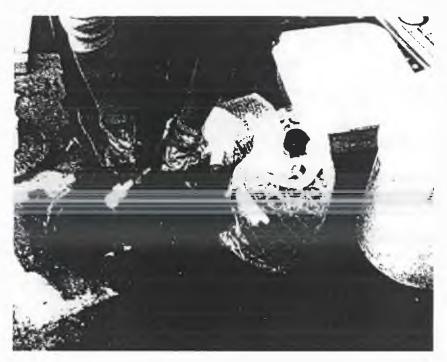
Date: May 6, 1991 Picture Taken By: Ben Farrell Direction Facing: N/A
Picture Description: Collecting volatile organic samples at monitoring well MW-D.

Photo No. 22



Date: May 6, 1991 Picture Taken By: Ben Farrell Direction Facing: N/A

Picture Description: VOA vial showing septum that fell out of vial cap.



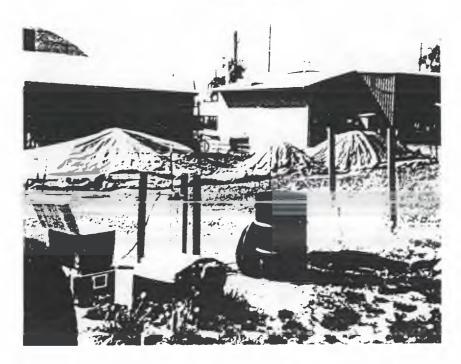
Date: May 6, 1991 Picture Taken By: Julie Howe Direction Facing: N/A

Picture Description: Filling pesticide samples at monitoring well MW-D. Note that the pump discharge tube is in contact with bottle opening.



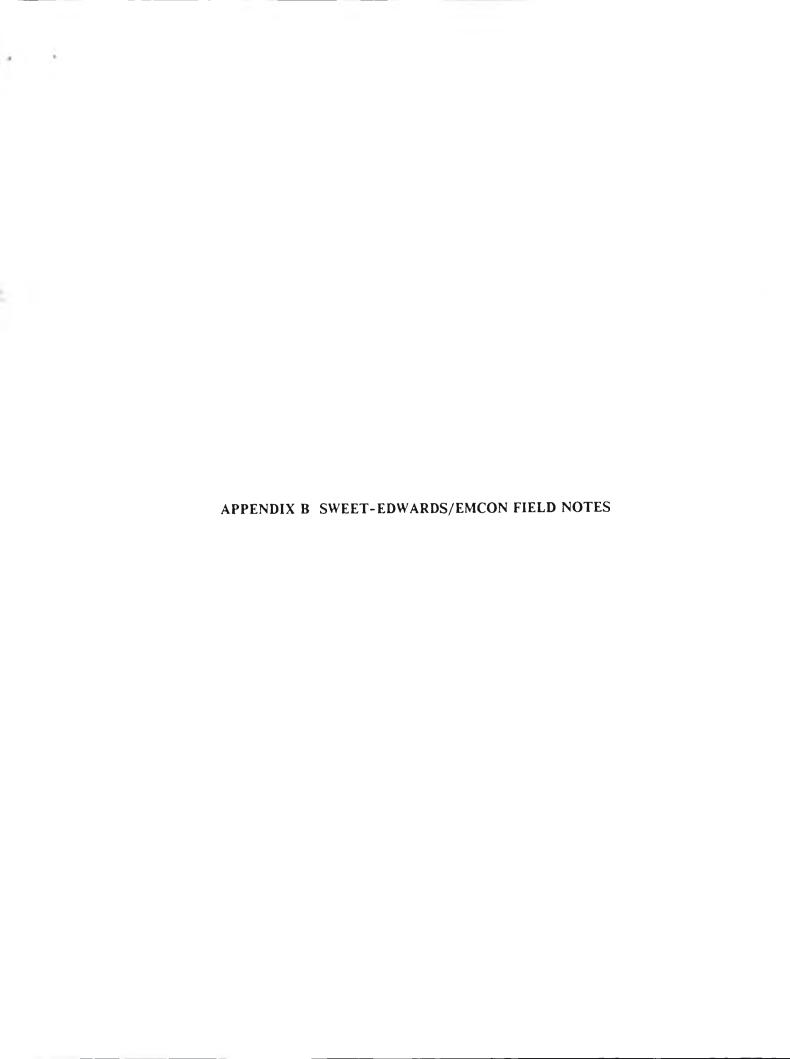
Photo No. 24

Date: May 6, 1991
Picture Taken By: Ben Farrell
Direction Facing: West
Picture Description: Collecting sample
for metals analysis at monitoring
well MW-C.



Date: May 6, 1991 Picture Taken By: Ben Farrell Direction Facing: West

Picture Description: Photograph of old drainfield area showing excavated soil piles. Monitoring well MW-E in foreground.





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Field Sampling Data

* Dupicate *

LOCATIONIADDRESS Vaking A	ROJECT NAME DULLATALL SCHOLAR 14/25-0103					Sample Designation 90042 - 0591-D.				
CLIENTICONTACT DOLG GAILER	China west	<u>5:53</u>	_	5/6/9/ 2011 Sun	A	0915				
2	J		AAAETUGI	main Dark	× 250	7 <u>E</u>				
(Nearest .01 ft.) EH	-	vation Date, Time			Method Used (M.Scope Number or Other) Slope Ind. cator #15227					
VELL EVACUATION: Gallons Pore Volumes	Method Used . ivell ive a more		Rinss Metho			Time 25 - 0915				
urlace Water Flow Speed	Measuremen	t Method		O	ate, Time _					
AMPLING: WEIL W. Eard										
Dole. Sample Time Method YOA =/4/91 8240 Resource 8150 4080 Method 7140 6010 7470-143	large Poli	Depth Taken (leet)	Fleid Filtered (yes,no) N At N	Frederica.	ced {yes.no	Sample: Cleaning Method Non-Phosphiane delergent west H2O ringe MeOH ringe Olstilled H2O ringe				
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OTES:						•				
Depth-461-35 = 11	-6-2									
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'						, !				
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otal # of Bottles				10,	ĉ.					
Harry St. West. Co.			Signature	- ideac	5 1-14	netand				



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LOCATIONIADORESSY	akima !	Jarice to		rel las	Sample Desi	onation	X41-0	1-67 591-61
PROJECT NAME	ri- Sama	ina	11 W25-01	03	Date, Time_	5/6/9/		1323
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YDROLOGY MEASUREMEN	TS:			•				
(Nearest .01 (t.) 32(R	Eleve	itlon	Date. TI		Slop	od Used IM-So	ope Numbe	r or Other
/ELL EVACUATION:				•				
	Volumes		thod Used		Rinse Metho	· .		. Time ধুর 10 কুতি
urtace Water Flow Speed	•		Measurement h	dethod		4 0	ste. 7ime	
AMPLING:								
Date, Bampie Time VOA 5//0 Descript	Method	Volume (mf) 21 42 22 4220	Container Type Class Angles Street	Depth Taken (feet)	Fleid Filtered (yea,no)	Preserva-	(yes,no)	Sample: Cleaning Method Non-Phosphruli, detergent wash H20 ringe MeOH ringe Distiffed H20 rinse
7 7.44 2 7.53 4 7.81	579 579 579 586 701	14.5 14.5 14.5	<u> </u>	Zecilion		1		
OTES:		•						
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			1	 				
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Foral # of Bottles:		,			1	· -,	, , , ,	11 1
1, 17					aignetur			SEA-400-0



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	ovation Oate, Time	• Method	Method Used (M-Scope Number or Other)				
· Xo.55	्राद्वित जरः	Slope Ind.	caty #15227				
WELL EVACUATION: Galigns Pore Volumes	Method Used Likel with 20 rd.	Rinse Method	1636 5/6/91 -				
Surface Water Flow Speed	. Measurement Me	thod	Date, Time				
Sample Time Method VON 16 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	Volume Container (mi) Type 2/47 Glass	Depth Field Taken Filtered (feet) (yes.no)	Preserva- Iced Cleani live (yes,no) Methr V. Non-Phos detergent H20 rir MeOH ri Olstilled rinse				
Pore Vol. Number pH Conductivity 7 7 7 970 3 7 68 997	150 Eh 200 150 Z	%ans					
NOTES:							
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,	Marader.stic Col	6 Y					
,	al avoder.stic Col	6 Y					
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PRG Mis cot Collect							



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Field Sampling Data

DCATION/ADDRE ROJECT NAME (L LIENT/CONTACT	sex dock	ر چي دار		* WES-21		Sample Desig		45-026	190	
YDROLOGY MEA		Elevi	ation	Sate, Tir	ne ,	Method Used (M-Scope Number or Other)				
ELL EVACUATIO	N: Pare Vo	lymes	Me	thod Used		Rinse Method		Date, Time		
urlaca Water Flow	Speed			Measurement M	lethod		04	ite. Time	•	
	ite, me N	feihod	Volume (ml)	Container Type /	Ocoln Taken (feet)	Field Filtered (yes,no)	Preserva- tive	(ced (yes,no)	Sample: Claaning Method Non-Phosphistic detergent wash H2O rinse MeOH rinse Distilled H2O rinse	
NOTES:	<u> </u>	nductivity								
2X-104-X	d (shift)	- 0,64	Cour	l into s	20 mb) 0	o online	Mica.	s 6.47	Carboy	
					۶۰۰		3			
Lab'	4					Signatur	Blue	2 2 XX	SEA-400-	



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CLIENTICOL	TACT Do	3 Geller	1. Hong "	1. a. 12 sec 1. W25. a 1. s. t.)		Date, Time		2801	(145	
(No	Y MEABUREME Pareal .01 (L.) 2.97		vetion	Date, The	5/6/9/	Method Used (M.Scope Number of Other) 5/JAC Ind. Control # 15227				
WELL EYAL Gallo		e Volumes		ethod Used	· · · · · · · · · · · · · · · · · · ·	Rinse Method		Date:	. Time - //45	
Surface Wate	or Flow Speed			Measurement N	iethod			ste, Time _		
Samole VIA Dese	Date, Time 5/6 /3	Method	Volume (ml)	Container Type 3 (A.3.) Griner (A.3.)	Depth Taken (feet)	Fleid Filtered (yes,no)	Preserva tive HC 1 r 12n e HN0n	(yes,no)	Sample: Cleaning Method Non-Phospholid detergent west H2O rinse MeOH rinse Distilled H2O rinse	
Pare Vol. Number 2 3 NOTES:	ph 6 33 7.15 7.32	Conductivity 106 1017 1034	Tem 15 15	2 2 2 3 3 4 4 5 5 6 5 6 6 6 6 6 6 6 6 6 6 6 6 6 6	uliuns					
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CLIENT/CONTACT 12-6	Samit on	25-01-0	Labora 3	Date, Time Weather	gnation 9/4 S/6/6	55°p	1/05
(Neerest .01 (1.)	Elevation	Oate, Tin		Meth	od Used (M-So	cope Number	or Other)
VELL EVACUATION: Gallons Pore Volume 4 5 3		Method Used [1] 11 12 7 4 7 d Measurement Method			d .	Date, Time 5/6/9/ 1045 - 10	
urface Water Flow Speed					D		
Date, Sample Time Met VCF 5/3/2	Volume (mn) (1) (1) (1) (1) (1) (2) (3) (3) (4) (5) (5) (6) (7) (7) (7) (7) (7) (7) (7) (7) (7) (7	Container Type Classic Direct Police Police Container Co	Depth Taken (feet)	Field Filtered (yes.no)	Preserve. Ital Haloz	iced (yes,no)	Sample: Cleaning Method Non-Phosphilic detergent wash H2O dinge MeOH rinsc Distilled H2O
Number pM Cond 7.80 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2	uctivity Tem	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	ileno				
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LOCATION/ADDRESS PROJECT NAME CLIENT/CONTACT	arthry Simpl		W25. 21		Well or Surface Sample Dealg Date, Time Weather	nation	10042-0 16191	
HYDROLOGY MEASUR (Negrest 01 to 25 7 4		vation	Date. Tic S/G/91 C		Metho	od Uaed (M-S	cope Number	r or Other)
Gallons	Pore Volumes		ethod Used		Rinse Method		16/4: 13	Time 20 - 1345 20 - 1613
Surface Water Flow Spi	eed		Measurement 4	lethod			ate, Time	
SAMPLING: Oste. Sample Time Signature Signature Oste. Oste.	Method	Volume (mil) 2 Y Y C. 1 SSD.	Container Type 31.45 24602.2	Depth Taken (feet)	Find Filtered iyes,not	Preserva- tive H_1	!ced {yes,no; 	Sample: Cleaning Method Non-Phosphiblic detergent wash H2O rinse MeOH rinse Distilled H2O
Pore Vot. Number pH 7 46 2 7.91	Conductivity (699 738 738 733	Tem	15° /6 / 32 -x 49					
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Total # of Bottlest	4/			J		Pd		\$EA-400-0



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CLIENTICONTACT DOLL GELLOC	Aar cultural P		Date, Time	iation 2	10-545	7
IYOROLOGY MEASUREMENTS: (Nearest .01 /tl.) El	levation Date 5/6/	17 0758	Method Stack	Used IM-Sci 100:44	Doe Number	or Other
VELL EVACUATION: Gaijons Pore Volumes	Method Used	d.	Rinse Method	. 41.ia	Date.	Time 1255
Surface Weter Flow Speed	. Measureme	ent Method		D:	ate, Time _	
Date. Sample Time Method 1/14 5/6/3 4/14/2 DEST	Volume Container (mi) Type (A) 40 40 40 40 40 40 40 40 40 40 40 40 40	(feet)	Field Filtered (yes,no)	Preserve- tive	(yes.no)	Sample: Cleaning Method Non-Phospitula detergent was: H2O ringe MeOH ringe Distilled H2O ringe
Pore Vol Number pH Conductivit 7.81 1030 7.71 22910 7.70 1046		Agallans				
METER W.	1 26 = 2					
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APPENDIX C PRC FIELD NOTES

5/6/12 7:30 Met with Beeky Hylland John North offices with well level Bigun mexicenests at MW-D \$20 Well pump wizard. canisters w. Nitroger MW-D Some rust on custing plastic with discolorations short (too short) services sent to Biogranics 12051 Indian Creek Ct Belt-ville Maryland 21) 905

purging commenced at 8:38

2" Inno Iruneter well

- shut pump off

- pumped directly into

plastic beaker. (rinsod out 12t)

BF:

20

- temp Faylor them 2173)
- Eh & DSPH3
- Ph

reject field sampling duta well construction report

shut pump off during measurements

2 gallors per well volume

Et /Ph met put on wet, dirty, well pump top

buckets - 7 gullons total

water placed in SSqullar

9:06 started sampling
- vorts
adjusted pressure during sampline
with her depace

- filled with ministury - not chatted for buildes

> 11 ITC. mately, tilled 2 4 liter bottles 2×1 liter polyethyler

> > 5 lide 16 pestroide sungles

slide 15 VOA samplish

Puplicate sample site

90042 - D. - 8591

Hony I mait/your
West D.

NO would

10:00 MWGsurveyed to top of easing martiel with pan

Ph/Eh ca(rbratal at start of day

为用

E.

total of 6 gellons pregal
MW-6 Ehlph
descritaminated probo with
distilled H20 between wells
4 well volumes parged

- Well lated 1035 - VOAS adjusted at too high

Order of sampling follows

11:50 MW-A.

VOA vals tend to have bubbles

prescruative purged from
Containers.

19 USA sampling
20 Simpling MW-A (inside casing)
alide 21+22 sampling post a MWA

Say

VOA & Pasticide sampling
MW-F

MW-F

new role of film

slats +-6 Soil Piles + MW-F

Mul-E deep sampling
piezometor

No sand pact

25

States 7-8, purmeter sampling.

MN-F

16:37 well secured

wells my ted with indelible

intr-cos not capped on MW-E.

MW-A not morted ends like with gad

MW-C. 1636

PH Eh Temp

8.51 008 17 Field blant

7.71 990 15°C 1

7.66 197 15°C 3 Bit

AP

1700 well secured
3 gallons puryed
421

BF

f - - - A

- T

YARL MON 05/06/91 0650 PRC arrived on site 0720 Becky Hylland, SEV arrived

Weather overiast 50'F

Newscrements in all wells using a Stope Indicator (2). electronic water sensor.

The we indicator was first rinsed with 1:1 HRDC methand (i distilled water (Culling Spring)) while on the reel just before the measurement is taken.

MW-D 34.73 for MW-G 37.68

when level is measured, to a mark on the owten caring

all stick up wells were locked with a Master And lock

SE = SWEET & dutirds

WATER LEVELS

51W

MW-D 34.73

MW-G 37.68

MW-B 37.18

MW-A 36.97

MW-F 36.75

MW-E 35.74

MW-C 36.55

The We indicator tape is palled to the successor an arrest on the cute

The well with a pount to open them

finished LUL measurements at 0815

wells will be purged with chalicated well with with purps (blackdon pumps)
Nirvogen is used as the compressed gas sorrce.

SE samples Will be six to Biospherics in Maryland 12051 Indian (reck Court Bellsville, MD, 20705

MW-D. 01 0838 began punging Total clepth. 46' 34.73 DTW. WC. Diameter !: SETTE WIN praye 6 Gal. punge wirte com + cocordens $(2) \quad (3)$ time 08461 08520858 gain pp 2 pH 6,32+7.03 7.35 7.33 923 719 cord 963 78mp 14° 0900 Pinished purging 0905 Chegan sampling 0930 finished sampling

MW-D SW #S 90042 - \$2591-01 90042 - \$30591-02(dup)

Problems Sampling due to
The Shortered liveth of
The Tubing from Miss
Well-inade it very
awkward for the Sample
to do field marnetes
She wad to keeps Stoping
The pirging process to

The tubing is langer to Some me had out it

She also lost a septa from a VIP VIAL & The efore control complete the sample set xxx: carey extra bottes.

MW-G

0254 organ purging mit 9

Total D 50' PTW 197.68 WC 12.32' CHILIMETER 2"

SW will june 6 sees (unless 2 intheld parameters are not with 102)

clean colones 30 oder

()

time 1000 1007 1011 1016
purged 2 gd 4 6 8
p.H 8.16 7.94 7.83 7.88
cond 579 598 686 701
14.5 14.5 14.5 14.5

1018 finished puging began sampling 1035 finished Sampling SW# 90042 0591 G1

Problems

flow rate was really to fast for collecting vox

septa - rinsed w/ punge water 4 used it

Recalibration of instruments

cond meter Set zero Set span cond STD 720 MMb

PH Set cal to 7.0 STD set Slope to 4.0 STD

MW-B

70tai D 47' DTW 37.18 WC 9.82 diameter 2"

SW well punge 4.5 gals for 3 (as Vs clear, colorless, no occar finished sampling 1602 SE Sample # 90042.0591-B1

PRC will not collect split samples from MW-BPlit

The flow raile was better for VOC sample

all pure water is cold sted in 55- Tal drums which are then primped out

Wery I got fine idea for 12 ha

began purging 1121

TD 42' DTW 36,97' WC 5.03'

SE WIll punge 3 gals for 3 in volumes Cliai, colorless, 10 dos

pursed 1 2 3
pH 6.33 7.15 7.32
cond 1106 1017 1034
T(r) 15 15 15

Stantial Samping 11:45
finished 12:35
700K extra volume form
ms/ms/s
and also took duplicate
(MW-K) at this well
sampling time 12:15

MINA (cont)

SE # 90042.0591-A1

Pump rate was
fast for VOAs

Ē

NOW SIRNY 70" MW-F

began purging 1235

punged 2 4 6 pH 7.82 7.71 7.60 Corld 1030 1038 1046 T 15 15 15

finished puring 1252 finished san pling 1313

SE Sample # 90042-0591-F1

sampling stoream was very acrated ".

Changed Nitrogen bottos began puigno 1320 finished 1610 began Sampling 1613 finished

Swe will stop until we can get a arm 1400 began priging agn, 1 1500

D D J time purged 3,1632 4853 PH 7.467.987 7.3999 Cond 699 738 733 T 15 15 15C Loranon: South of mws E, F

1340 at 17 Juls pursed no field measurements have been taken because Becky is calling H because she is running out of space in the purguent, culectionarums

TD 128"

DTW 35,74"

WC 92.26"

Clin J"

			Ageny units products	
	0.0	washington and the		
*11	to see market estimation to the see	. The Anticophical Ministry of the Constant of	all Mart Bare 1444 Andréin (1888)	
and the second s				to Kingletone
			124.04	
	12141		204 - 14 14 2 14 2 4 CC	
		•		
		j		
	APPENDIX D OPER	ATION AND MAINTENAN	CE FIELD INSPECTION	N CHECKLIST
				-
	-			

4. Observe the owner/operator's staff as they collect ground-water samples at several wells. Complete the following table for each well (Note: revise or add to the table if permit conditions dictate a different requirement the owner/operator must follow):

Position/Title	Name	Sampling Experience (years and type)				
Geologist	Beeky Hylland	2/yr. of environmental				

Well Identification Number Mw-D (Also applies to unst other wells)	Y/N	Photograph Taken Y/N
Did the sampling crew measure static water levels in the well and well depths prior to the sampling event?	N	N
Did the sampling crew use a steel tape or electronic device totake depth measurements?	Y	7
Did the sampling crew record depths to +/- 0.01 feet?	4	4
Did the sampling crew follow these procedures: 1. remove locking and protective cap; 2. sample the air in the well head for organic vapors; 3. determine the static water level; and 4. lower an interface probe into the well to detect immuscible layers.	γ	\mathcal{N}
If immiscible samples were collected, were they collected prior to well purging?	WA	_
Did the sampling crew evacuate low yielding wells to dryness prior to	N	
Did sampling crew evacuate high yielding wells so that at least three casing volumes were removed?	7	7
Did the sampling crew collect the purge water for storage and analysis or for shipment off-site to a RCRA treatment facility?	N	N
Were sampling devices constructed of fluorocarbon resins or stainless	NA	_

Well Identification Number	Y/N	Photograph Taken Y/N
if the sampling crew used dedicated samplers, did they disassemble and thoroughly clean the devices between samples?	NA	- 1
If samples are collected for organic analyses, did the cleaning procedure include the following steps: I non phosphate detergent wash Lap water runse distilled/detonized water runse acetone runse pesticide-grade hexane runse?	N	y
If samples are collected for inorganic analyses, does the cleaning procedure include the following steps: i. dilute acid rinse (HNO ₃ or HCL) 2. distilled/de-ionized water rinse?	\sim	n
Did the sampling crew take trip blanks, field blanks and equipment blanks? Yes Yes Wo	7	\sim
If the sampling crew used bailers, were they bottom valve bailers?	NA	_
If the sampling crew used bailers, was "teflon" coated wire, single strand stainless steel wire or monofilament used to raise and lower the bailer?	NA-	_
If the sampling crew used bailers, did they lower the bailer slowly to the well?	NA	-
If the sampling crew used bailers, were the bailer contents transferred to the sample container to minimize agitation and aeration?	NA	_
Did the sampling crew take care to avoid placing clean sampling equipment, hoses, and lines on the ground or other contaminated surfaces prior to insertion in the well?	7	N
f the sampling crew used dedicated bladder pumps: Was the compressed gas from an oilless compressor certified quality commercial compressed gas cylinder? If not, was a suitable oil removal purification system installed and maintained?	7	N
Vas the bladder pump controller capable of throttling the bladder pump ischarge flow to 100 mi/min or less for continuous periods of at least 0-30 seconds without restricting liquid discharge?	not Known	N

Well Identification Number Mw-D	Y/N	Photograph Taken Y/N
Were samples taken from the bladder pump discharge tube, and not from any purge device discharge tube?	.7	7
Was the bladder pump discharge flow checked for the presence of gas bubbles before each sample collection, as a test for bladder integrity?	N	N
Was bladder pump flow performance monitored regularly for dropoff in flow rate and discharge volume per cycle?	y	rv
Was the bladder pump incorporated in a combination sample-purge pump design which can expose the bladder pump interior and discharge tubing to the pump drive gs? If so, were operating procedures established and followed to prevent at all times the entry of drive gas into the sample flow or into the bladder pump interior?	N	N
Did the sampling crew collect and containerize samples in the order of the volatilization sensitivity of the parameters?	Y	Y
Did the sampling crew measure the following parameters in the field: pH, temperature, specific conductane?	Y	4
Did the sampling crew sample background wells before sampling downgradient wells?	Y	4
Did the sampling crew use fluorocarbon resin or polyethylene containers with polypropylene caps for samples requiring metals analysis?	7	4
Did the sampling crew use glass bottles with fluorocarbon resin- lined caps for samples requiring metals analysis?	N	N
If metals were the analytes of concern, did the sampling crew use containers cleaned with nonphosphate detergent and water, and insed with nitric acid, tap water, hydrochloric acid, tap water and finally Type II water? (leanel of lab	un trong	
f organics were the analytes of concern, did the sampling crew use containers cleaned with nonphosphate detergent, rinsed with tap vater, distilled water, acetone, and finally pesticide quality hexane?	untnow	-
oid the sampling crew filter samples requiring analysis for organics?	No	_

2. Visually inspect each well and piezometer and complete the table below (one line entry for each well or piezometer):

					minor 1 451			
Well/ Piezometer	Survey Mark Present?	Standing or Ponded Water?	Evidence of Collision Damage?	Evidence of Frost Heaving?	Evidence of Casing Degradation?	Lock in Place?	Evidence of Well Subsidence?	Photograph Taken?
mw-p	Tes	Standing water Finall wells.	W_{o}	No	minor rust	flush mount- bolted	No	185
wm-C	745	No starting water outside	N_{o}	Wo	No	Yes	Wo	Ya,
Mw-B	Yes	otcasing	No	No	No	Yes	14	Yes
MW-P	Tes		\mathcal{N}_{o}	N	No	404	No	Yes
Mw-F	765		n,	N	No	Yes	nz	Yes
MW-E	Ye5		No	N	N_{o}	Yez	No	Yes
MW-C	Yes		Nb	No	N6	Ye5	No	705

APPENDIX E YARL ANALYTICAL DATA SUMMARY

BIOSPHERICS INCORPORATED

CLIENT: Hong West

DATE COLLECTED: May 6, 1991

DATE RECEIVED: May 9, 1991

MATRIX: Water

LAB I.D.: 91-05-0916

Analytical Methodology/Sample Chronicle

Parameter	Method	Date Extracted	Date Analyzed
Metals	EPA 6010/7000's	5/15/91	5/14-29/91
Mercury	EPA 7470	5/21/91	5/21/91
Pesticides/PCB's	EPA 8080	5/10/91	5/13-14/91
Organophosphorus Pesticides	EPA 8140	5/13/91	5/16/91
Volatile Organics	EPA 8240		5/18/91
Herbicides	EPA 8150	5/13/91	5/15/91

Non-conformance Summary

Pesticides/ PCB's

Endrin yielded low recoveries (<56%) in the spike blank, matrix spike, and matrix spike duplicate. Surrogate recoveries were acceptable in all samples. Since holding time expired, samples were not reextracted. Lindane recoveries in the matrix spike and matrix spike duplicate were below acceptable levels (<56%). However, the spike blank was acceptable and the data was released with confidence.

Organophosphorus Pesticides

Due to a spiking error TEPP recovery was low (<40%) in all quality control samples. The TEPP analyses cannot be reported with confidence.

SUMMARY OF MAY 6, 1991 MONITORING YAKIMA AGRICULTURAL RESEARCH LABORATORY

Split Sample Collection: Split samples taken from MW-G, MW-D, MW-A, MW-F and MW-E.

Summary of Results (units are ug/L)

ND - denotes non-detected or below quantitation limit

F , Et Z

WELL ID	volatile org. 8240	TCL Metals	Pest/PCB 8080	Insect/Herb 8140/8150
MW-G	ND	Ca 74,200 Mg 46,600 K 3,540 Nn 64,300 V 75.8	ND	ND
MW-D	ND	Ba 57.3 Ca 73,500 Mg 45,700 K 3,210 Na 68,000 V 74.9 Zn 27.0	ND	ND
MW-A	ND	Ca 76,000 Pb 3.5 Mg 43,200 K 4,030 Na 62,500 V 57.2 Zn 20.3	ND	ND
MW-F	ND	Ca 78,200 Mg 45,400 K 3,610 Na 61,800 V 71.9 Zn 39.7	ND	ND
MW-E	ND	Ca 59,300 Mg 28,600 K 4,820 Na 30,100	ND	ND

APPENDIX F PRC ANALYTICAL DATA SUMMARY

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TABLE 1 INORGANIC ANALYTICAL RESULTS, YAKIMA AGRICULTURE RESEARCH LABORATORY $(\mu g/L)$

Sample Location:	MW-D T D	MW-A T D	MW-A T D	MW-E T D	MW-F T D	T D
Contract Laborate Program Number:	ory	MJF144 MJF145	MJF146 MJR147	MJF413 MJF414	NUF148 MUF149	MJF125 MJF143
Aluminum	14.0U 14.0U	14.0U 14.0U	28.8 15.1	14.0U 14.0U	17.2 31.4	14.0U 19.5
Arsenic	7.2 7.2	5.2 5.6	4.7 4.9	3.0 2.8	5.4 6.1	6.3 6.9
Barium	21.6 21.1	20.5 25.2	20.0 20.6	24.2 24.6	13.0 17.7	18.3 18.2
Calcium ,	71,400.0 74,500.0	79,800.0 78,500.0	79,100.0 78,600.0	57,900.0 55,100.0	74.100.0 76,400.0	75,700.0 71,400.0
Copper	4.3 4.0U	4.0U 4.0U	4.0U 4.0U	4.0U 4.0U	4.0U 4.0U	4.0U 4.0U
Iron	34.1 8.0U	12.1 8.0U	20.1 13.2	10.4 8.0U	11.5 8.0U	8.0U 8.0U
Lead	5.5 2.7	2.7 3.4	17.2 3.0	1.0U 3.0	2.9 4.1	3.0 3.9
Magnesium	44,300.0 45,600.0	44,000.0 43,600.0	43,100.0 42,500.0	27,200.0 26,800.0	43,400.0 44,600.0	46,700.0 44,900.0
Potassium	3,290.0 3,320.0	4,190.0 4,190.0	4,070.0 4,020.0	4,790.0 4,800.0	3,410.0 3,550.0	3,550.0 3,570.0
Sodium	67,200.0 68,600.0	65,800.0 65,400.0	63,100.0 62,000.0	30,100.0 30,500.0	59,500.0 61,700.0	67,300.0 65,600.0
Vanadium	66.6 68.0	52.2 52.2	50.8 54.4	26.5 27.1	59.1 60.5	67.7 65.0
Zinc	11.3 8.7	7.0U 7.0U	7.0U 21.7	7.0U 7.0U	9.1 7.0U	7.0U 7.0U

Notes:

Metals not listed were not found in any samples. T = Total metals.

D = Dissolved metals (sample filtered in the field).

Qualifier: U = Not detected; listed value is the sample detection limit.

TABLE 2 ORGANIC ANALYTICAL RESULTS, YAKIMA AGRICULTURE RESEARCH LABORATORY $(\mu g/L)$

Sample Location:	Trip Blank	MW-D	MW-A	MW-A	MW-E	MW-F	MW-G
Contract Laboratory Program Number:	JG889	JG879	JG885	JG886	JG888	JG887	JG884
Volatile Organics							
Methylene chloride	2.J	5U	5U	5U	5U	5U	5U
Acetone	10U	10U	10U	10U	4J	5J	10U
Chloroform	5U	5U	2.J	1J	1J	2.J	3Ј
Tetrachloroethene	5U	3J	5U	5U	5U	1,3	5U
Pesticides and PCBs							
Endosulfan sulfate	NS	0.10U -	0.20	0.17	0.10	0.10	0.10

Notes:

Chemicals not listed were not found in any samples (including all target analytes in the organophosphorus pesticide and chlorinated herbicide assays).

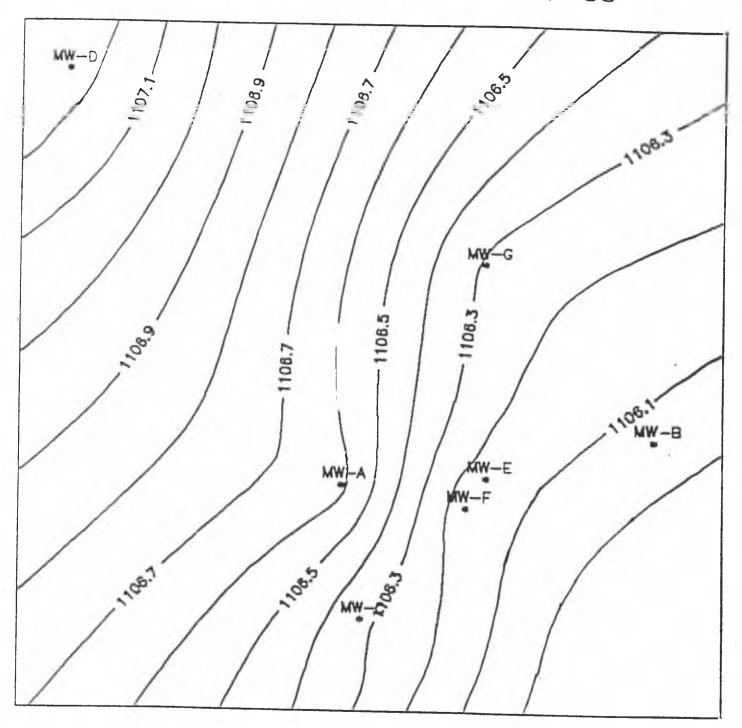
Qualifiers:

U = Not detected; listed value is the contract-required quantitation limit (CRQL).
 J = Estimated value; most commonly the chemical was found at a concentration less than the CRQL.

NS = No sample.

APPENDIX G POTENTIOMETRIC SURFACE MAPS FOR THE YARL SITE

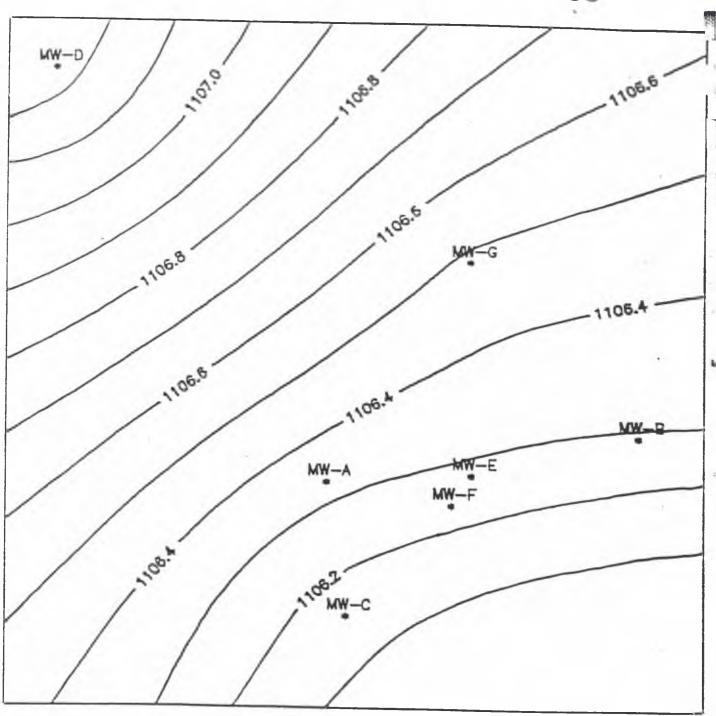
YARL GROUNDWATER LEVEL 8-7-90



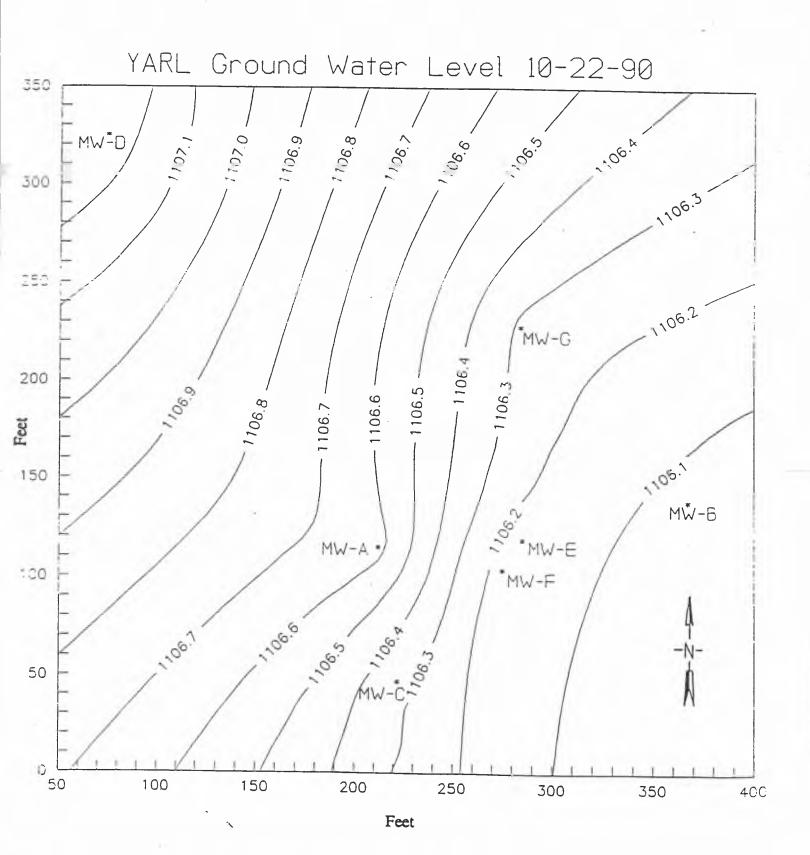
SCALE 1 inch = 50 FEET

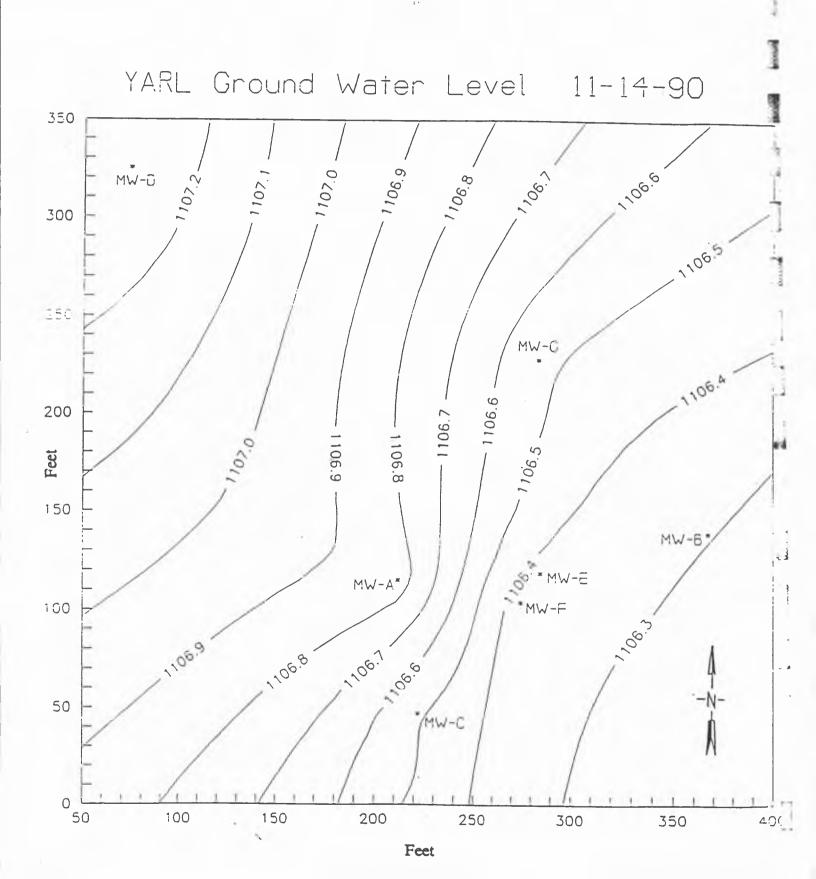
YARL GROUNDWATER LEVEL 9-4-90

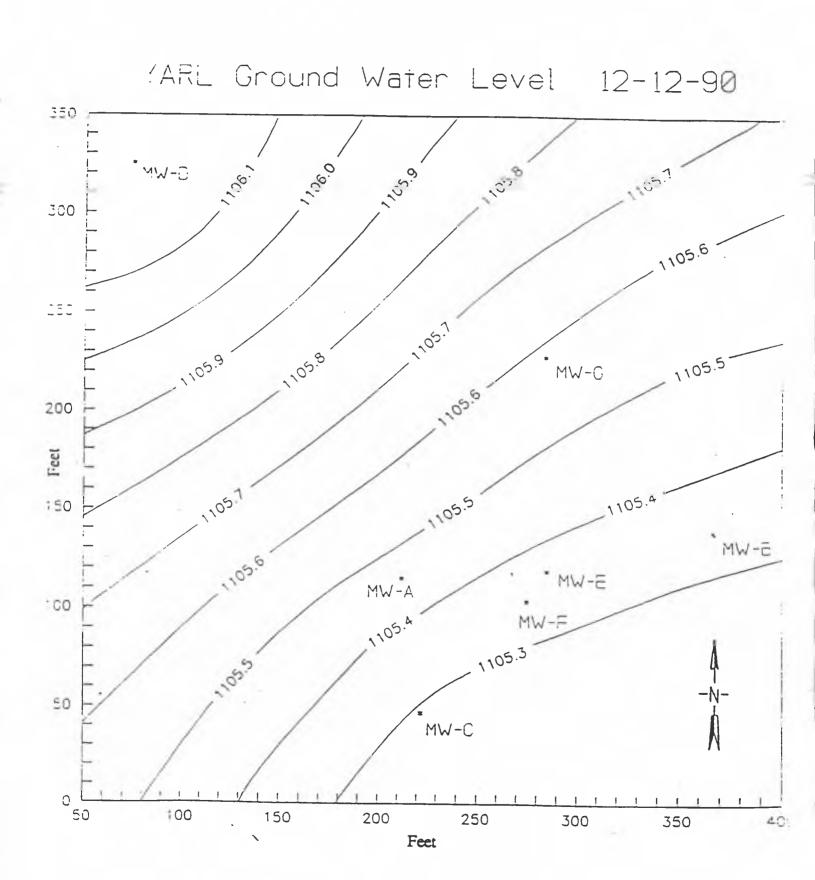
5 12

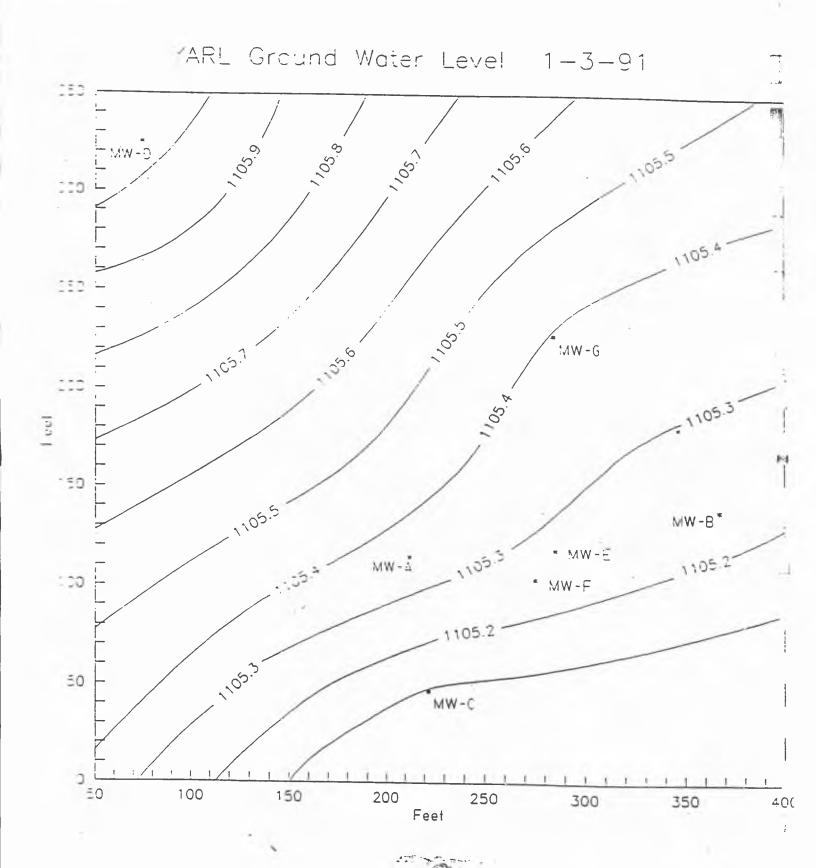


SCALE 1 inch = 50 FEET

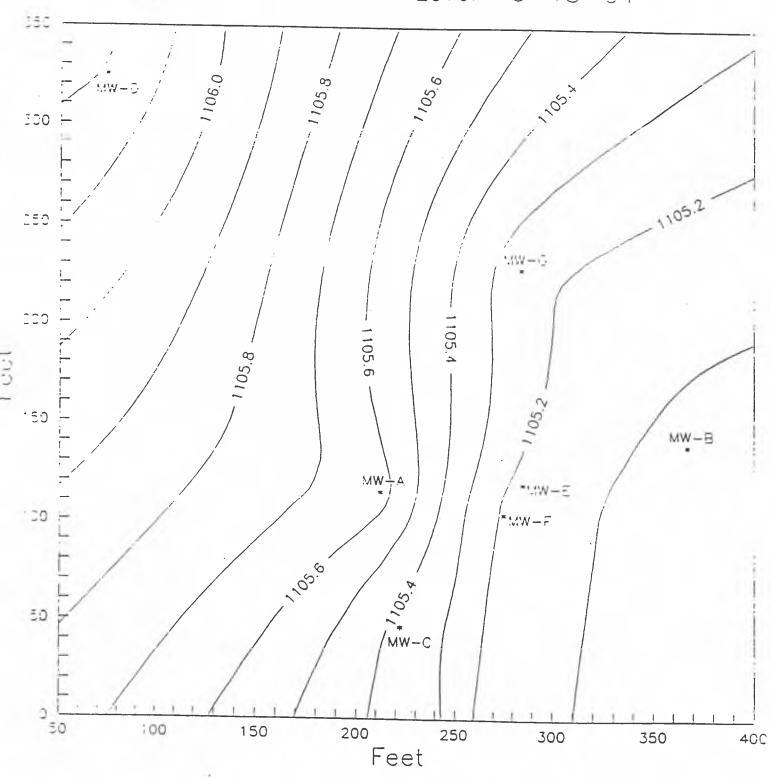


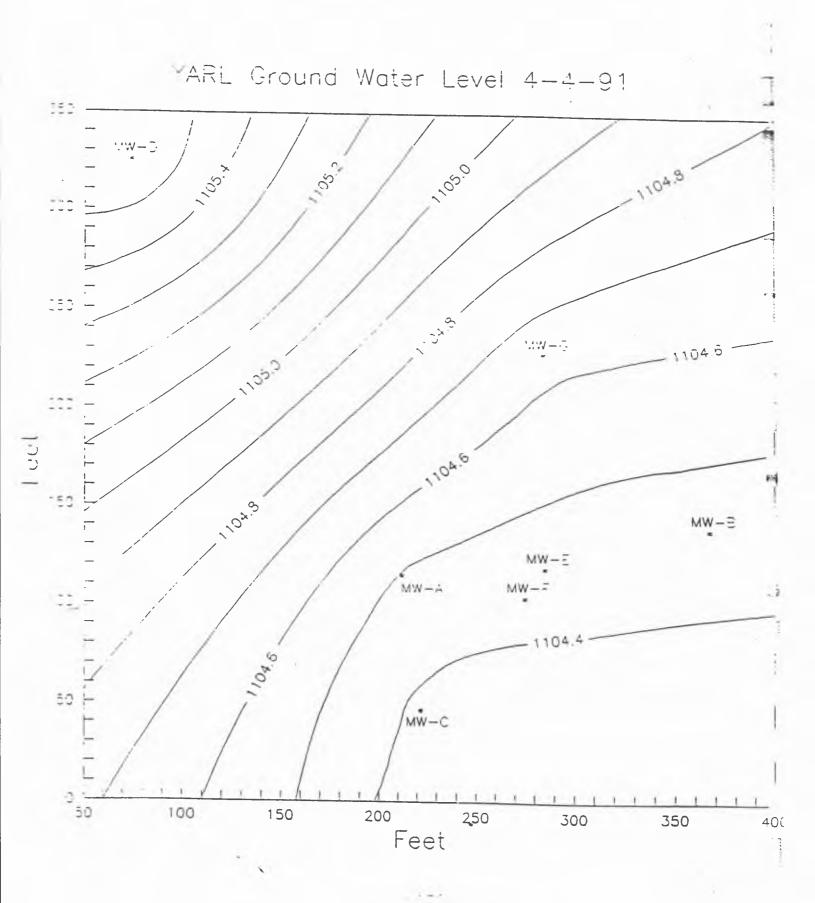


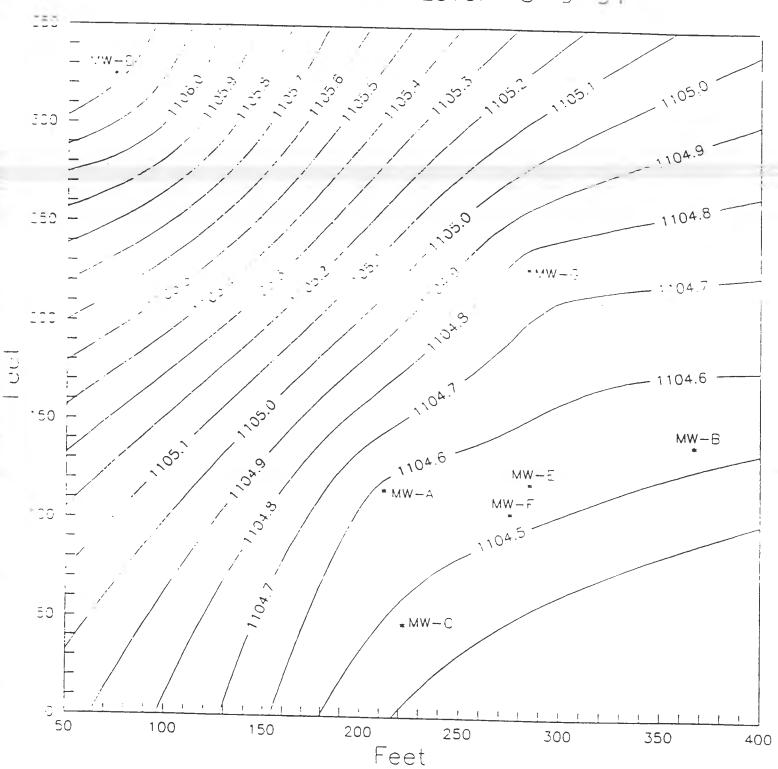




ARL Ground Water Level 3-13-91







ARL Ground Water Level 6-13-91 105.3 ,105.2]]] -1105.1 250 -1105.0 .,105.0 1104.9 ·50 -1104.9 MW-B .00.0 MW-EMW-A MW-F 11047 ED * MW-C 100 150 200 250 300 350

Feet

40

